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(21) International Application Number: PCT/EPS (22) International Filing Date: 18 February 1999 (1998) (30) Priority Data: Mi98A000356 24 February 1998 (24.02.98) (71) Applicant (for all designated States except US): 25 S.P.A. [IT/IT]; Galleria del Corso, 2, I-20122 Miller (1998) (72) Inventors; and (1998) (73) Inventors/Applicants (for US only): BARTORELLI [IT/IT]; Via G. D'Arezzo, 6, I-20145 Milano (IT) Cesare [IT/IT]; Galleria del Corso, 2, I-20122 Miller (1998) (74) Agent: MINOJA, Fabrizio; Bianchetti Bracco Minoja (1998) (75) Rossini, 8, I-20122 Milano (IT).	) I ZETES: ano (IT , Alber , SANT lano (IT	(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, IP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  Published  With international search report.  Before the expiration of the time limit for amending the

(54) Title: ORAL COMPOSITIONS AT LOW DOSAGE OF CYTOTOXIC PROTEINS

#### (57) Abstract

Pharmaceutical compositions for the oral and sublingual administration containing proteins extractable from mammalian liver. Such proteins include the proteins marked with UK101 and UK114 and described in WO 92/10197 and WO 96/02567 as well as ubiquitin, contained in the UK101 extract.

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# ORAL COMPOSITIONS AT LOW DOSAGE OF CYTOTOXIC PROTEINS

The present invention relates to pharmaceutical compositions for oral and sublingual administration containing proteins extractable from mammalian liver.

Such proteins include the proteins referred to as UK101 and UK114 and described in WO 92/10197 and WO 96/02567 as well as ubiquitin, contained in the UK101 extract.

It has been observed that the subcutaneous administration of UK101 and UK114 induces a clear cytotoxicity in serum of both healthy subjects and in tumor carriers subjects.

The main responsible for this effect is the 14Kd protein (UK114) contained in the protein extract UK101. The UK114 amino acidic sequence has been described in FEBS Let. 393, 147-150, 1996.

At present, clinical experimentations to verify the UK101 and UK114 therapeutic efficacy are in progress. To this order, patients suffering from colon and breast carcinoma are treated with UK101 and UK114 subcutaneous injections, at dosages ranging from 1 to 10 mg/week.

The protein nature of the active principle obviously forces the parenteral route

In fact, at present it is not known the possibility to administer proteins orally, due to their high metabolic instability.

To avert this inconvenience, it has been suggested several answers such as the use of suitable carriers or the encapsulation in liposomes, but until now the results have been unfavourable.

Now it has been found that it is possible to administer UK114 and UK101 or ubiquitin orally at low dosages, preferably for sublingual

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administration, thus inducting a seric cytotoxicity comparable to or higher than that obtainable for subcutaneous administration.

The oral/sublingual route also shows clear advantages in practicality and safety terms.

The invention therefore provides pharmaceutical compositions for UK101 and UK114 oral and/or sublingual administration.

Suitable administration forms include, for example, aqueous suspensions to administer in drops, granules or sublingual tablets by a quickly disintegration, effervescent or chewable tablets or its equivalent forms.

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The compositions of the invention can be prepared using conventional techniques and excipients, widely known in the pharmaceutical field.

The UK101 and UK114 unitary dosages can range from  $10 \times 10^{-4}$  to  $10 \times 10^{-15}$  g.

In the case of solutions to administrate in drops, the concentration of the active principle can range from 10<sup>-5</sup> to 10<sup>-10</sup> M. The administration of 10-15 drops a day proved to be sufficient to induce cytotoxicity in the patient serum, which can be evidenced on carcinoma cells Jurkat and Kato III according to standard protocols.

### 20 EXAMPLE

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17 patients suffering from tumors in advanced phase (8 sarcoma, 4 breast carcinoma, 2 pancreas carcinoma and 3 colon carcinoma) have been treated with 5-20 drops/day of an 1% hydroalcoholic solution of ethanol in a UK101 concentration of 10<sup>-6</sup> M.

The treatment continued for 30 consecutive days, involved in 70% of the cases an improvement in the subjective conditions of the patients, particularly a decrease in the painful symptomatology, a decrease in the tumor mass in 20% of the cases, joined to a cytotoxicity visible in the

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patients serum on cell lines Jurkat and Kato III.

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### **CLAIMS**

- 1 Oral pharmaceutical compositions containing as active principle a protein selected from ubiquitin, UK114 and UK101.
- 5 2 Composition according to claim 1, suitable for the sublingual administration.
  - 3 Compositions according to claim 2, in the form of drops, granules or sublingual tablets.

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4 Compositions according to any one of the previous claims, 10 containing from 10<sup>-4</sup> to 10<sup>-15</sup> g of UK101, UK114 and ubiquitin for unitary.

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### INTERNATIONAL SEARCH REPORT

Int Itional Application No PCT/EP 99/01068

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C. DOCUM	IENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the re	elevant passages	Relevant to claim No.
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Fun	ther documents are listed in the continuation of box C.	Patent family members are listed	l in annex.
"A" docum consi "E" earlier filing "L" docum which citatic "O" docum other "P" docum later t	ent which may throw doubts on priority claim(s) or is cited to establish the publication date of another on or other special reason (as specified) nent referring to an oral disclosure, use, exhibition or means ent published prior to the international filing date but than the priority date claimed	"T" later document published after the interpretation or priority date and not in conflict with cited to understand the principle or the invention.  "X" document of particular relevance; the cannot be considered novel or cannot involve an inventive step when the document of particular relevance; the cannot be considered to involve an indocument is combined with one or ments, such combination being obvious in the art.  "&" document member of the same patent	the application but secry underlying the claimed invention to considered to coument is taken alone claimed invention eventive step when the ore other such docurus to a person skilled
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